



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY
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HEALTH EFFECTS DIVISION
SCIENTIFIC DATA REVIEWS
EPA SERIES 361

OFFICE OF
PREVENTION, PESTICIDES, AND
TOXIC SUBSTANCES

MEMORANDUM

DATE: 14-APR-1999

SUBJECT: **Pyridate - Acute and Chronic Dietary Exposure Analysis.** PP#: 99MT0012. Chemical#: 128834. DP Barcode: D254602. Case #: 291638. Submission #: S558560.

FROM: Jennifer E. Rowell, Chemist *Jennifer E. Rowell*
Registration Action Branch 1
Health Effects Division (7509C)

THROUGH: Melba Morrow, D.V.M., Branch Senior Scientist *Melba Morrow*
Registration Action Branch 1
Health Effects Division (7509C)

TO: George F. Kramer, Ph.D., Chemist
Registration Action Branch 1
Health Effects Division (7509C)

Action Requested

Provide an estimate of the dietary exposure and associated risk for pyridate resulting from existing tolerances and proposed tolerance level for residues on mint (PP# 99MT0012). The petitioner proposes the establishment of time-limited tolerances for the residues of pyridate in or on the following raw agricultural commodity (RAC):

Mint - 0.3 ppm

Tolerances have been established for the residues of pyridate (40 CFR §180.462) in or on cabbage at 0.03 ppm; corn, fodder at 0.03 ppm; corn, forage at 0.03 ppm; corn, grain at 0.03 ppm; corn, silage at 0.03 ppm; and peanut, nutmeat at 0.03 ppm. A Section 18 time-limited tolerance (expiration/revocation date 12/31/98) has been established for pyridate in or on chickpeas at 0.1 ppm.

Executive Summary

Acute and chronic dietary exposure analyses for pyridate were performed using the Dietary Exposure Evaluation Model (DEEM™). Acute and chronic dietary exposure analyses were performed using tolerance level residues and 100% crop treated (CT) information for all commodities. All dietary risk estimates are below the Agency's level of concern for the U.S. population and sub-populations (including infants and children).

Toxicological Endpoints

On October 21, 1997, the Health Effects Division's Hazard Identification Review Committee (HIARC) met to evaluate the toxicology data base of pyridate with special reference to the reproductive, developmental and neurotoxicity data. These data were re-reviewed specifically to address the sensitivity of infants and children from exposure to pyridate as required by the Food Quality Protection Act (FQPA). In addition, the Committee also re-assessed the doses and endpoints selected for acute dietary, chronic dietary (RfD) as well as occupational and residential exposure risk assessments (Memo, J. Rowland 11/3/97). A summary of the toxicological endpoints chosen by HIARC is listed in Table 1.

Acute

For the acute analysis, the HIARC selected an RfD of 0.2 mg/kg/day (NOAEL = 20.0) based on ataxia and emesis observed within 1-3 hours dosing beginning on the first day in a 90-day dog feeding study at a LOAEL of 60 mg/kg/day. An uncertainty factor (UF) of 100x was applied to account for both inter-species extrapolation [10x] and intra-species variability [10x] (Memo, J. Rowland 11/3/97).

Chronic

For the chronic analysis, the HIARC selected an RfD of 0.11 mg/kg/day (NOAEL = 10.8) based on decreased body weight gain in males observed in a 2-year feeding study at a LOAEL of 67.5 mg/kg/day. An uncertainty factor (UF) of 100x was applied to account for both inter-species extrapolation [10x] and intra-species variability [10x] (Memo, J. Rowland 11/3/97).

Table 1. Summary of Toxicological Endpoint Selection

EXPOSURE SCENARIO	DOSE (mg/kg/day)	ENDPOINT	STUDY
Acute Dietary	NOAEL = 20 UF = 100	Clinical signs indicative of neurotoxicity characterized as ataxia and emesis were observed within 1-3 hours post-dosing on the first day and persisted for duration of study. LOAEL = 60 mg/kg/day.	90-Day Feeding Study - Dog
		Acute RfD = 0.20	
Chronic Dietary	NOAEL = 10.8 UF = 100	Based on decreased pup weight gain (at post natal day 14 and 21 in the first litters of both generations) at the LOAEL of 67.5 mg/kg/day.	Chronic Toxicity/Carcinogenicity Study - Rat
		Chronic RfD = 0.11	
Short-Term (Dermal)	NOAEL = 20	See Acute Dietary.	90-Day Feeding Study - Dog
Intermediate-Term (Dermal)	NOAEL = 20	See Acute Dietary.	90-Day Feeding Study - Dog
Long-Term (Dermal)	NOAEL = 10.8	See Chronic Dietary.	Chronic Toxicity/Carcinogenicity Study - Rat
Short Term (Inhalation)	NOAEL = 20	See Acute Dietary.	90-Day Feeding- Dog
Intermediate Term (Inhalation)	NOAEL = 20	See Acute Dietary.	90-Day Feeding- Dog
Long Term (Inhalation)	NOAEL = 10.8	See Chronic Dietary.	Chronic Toxicity/Carcinogenicity- Rat

FQPA Recommendation

On April 5, 1999, the FQPA Safety Factor Committee met and determined that the 10x factor to account for enhanced sensitivity of infants and children should be removed. **Please note: the decisions made at this meeting for this chemical are applicable only to this Section 18 request (Memo, B. Tarplee 4/8/99).**

The Population Adjusted Dose (PAD) is a modification of the acute or chronic RfD to accommodate the FQPA Safety Factor. The PAD is equal to the acute or chronic RfD divided by the FQPA Safety Factor. **Since the HED FQPA SFC determined to remove the 10x safety factor, the RfD is identical to the PAD.**

Cancer

Pyridate has not been to the Cancer Peer Review Committee. However, the DERs for the mouse and rat oncogenicity studies indicate that pyridate was negative in both species for carcinogenic effects (Memo, A. Kocialski, et. al. 7/11/97).

Residue Information

Tolerances for pyridate are published in 40 CFR §180.462. For the acute and chronic analyses, tolerance level residues and 100% CT information were used for all commodities. A summary of the residue information used in the acute and chronic analyses is attached (Attachment 1).

Results/Discussion

The Dietary Exposure Evaluation Model (DEEM™) analysis evaluated the individual food consumption as reported by respondents in the USDA 1989-91 Nationwide Continuing Surveys of Food Intake by Individuals (CSFII) and accumulated exposure to the chemical for each commodity.

Acute Dietary Exposure Analysis

The acute dietary exposure analysis estimates the distribution of single-day exposures for the U.S. population and certain subgroups and accumulates exposure to the chemical for each commodity. Each analysis assumes uniform distribution of pyridate for the commodities on which pyridate is used.

A Tier 1 acute analysis was performed using published and proposed tolerance levels and 100% CT information for all commodities (Attachment 1). For acute dietary risk, HED's level of concern is >100% PAD. Dietary exposures and associated acute risk at the 95th percentile are shown in Table 2. The other subgroups included in Table 2 represent the highest dietary exposures for their respective subgroups (i.e., children, females, and the other general population subgroups higher than U.S. population). A full listing of dietary exposures is attached (Attachment 2).

Table 2- Summary of Results of Acute DEEM Analysis for Pyridate at the 95th Percentile.

Subgroups	Exposure (mg/kg/day)	% PAD
U.S. Population (48 states)	0.000139	0.1
Non-Hispanic Blacks	0.000159	0.1
Non-nursing Infants (<1 yr)	0.000278	0.1
Children (1-6 years)	0.000268	0.1
Females (13+/nursing)	0.000124	0.1

The results of the acute analyses indicate that the acute dietary risk associated with the existing and proposed uses of pyridate is well below the Agency's current level of concern.

Chronic Dietary Exposure Analysis

The chronic dietary exposure analysis used mean consumption (3-day average) data. A Tier 1 analysis was performed using published and proposed tolerance level residues and 100% CT information for all commodities. For chronic dietary risk, HED's level of concern is 100% PAD. Dietary exposures for the General Population and other subgroups are presented in Table 3. The other subgroups included in Table 3 represent the highest dietary exposures for their respective subgroups (i.e., children, females, and the other general population subgroups higher than U.S. population). A full listing of chronic dietary exposures is attached (Attachment 3).

Table 3. Summary of Results from Chronic DEEM Analysis of Pyridate.

Subgroups	Exposure (mg/kg/day)	% PAD
U.S. Population (48 states)	0.000044	0.0
Non-Hispanic Blacks	0.000050	0.0
Non-nursing Infants	0.000121	0.1
Females 13-19 (not preg or nursing)	0.000043	0.0
Males 13-19 yrs	0.000054	0.0

The results of the chronic analysis indicate that the chronic dietary risk associated with the existing and proposed uses of pyridate is well below the Agency's current level of concern.

Cancer Dietary Exposure Analysis

Pyridate has not been to the Cancer Peer Review Committee. However, the DERs for the mouse and rat oncogenicity studies indicate that pyridate was negative in both species for carcinogenic effects (Memo, A. Kocalski, et. al. 7/11/97). Therefore, no cancer dietary exposure analysis was performed.

Conclusions

The acute analysis was performed using published and proposed tolerance levels and 100% CT information for all commodities (Attachment 1). The acute %PADs were <100%, and the highest was 0.1% for non-nursing infants (<1 year). For acute dietary risk, HED's level of concern is 100% PAD. The results of the acute analysis indicate that the acute dietary risk associated with the existing and proposed uses of pyridate is well below the Agency's current level of concern.

For the chronic analysis, a Tier I analysis was performed using published and proposed tolerance level residues and 100% CT information for all commodities. The %PADs for all subgroups were <100%, and the highest was 0.1% for non-nursing infants. The results of the chronic analysis indicate that the chronic dietary risk associated with the existing and proposed uses of pyridate is well below the Agency's current level of concern.

Pyridate has not been to the Cancer Peer Review Committee. However, the DERs for the mouse and rat oncogenicity studies indicate that pyridate was negative in both species for carcinogenic effects (Memo, A. Kocalski, et. al. 7/11/97. Therefore, no cancer dietary exposure analysis was performed.

- Attachment 1: Pyridate Residue File for Acute and Chronic DEEM™ Analyses.
- Attachment 2: Pyridate Acute DEEM™ Analysis (J. Rowell, 02-APR-1999).
- Attachment 3: Pyridate Chronic DEEM™ Analysis (J. Rowell, 02-APR-1999).

cc (w/attachments): J.Rowell (RAB1); M.Sahafeyen (CEB1); PP# 99MT0012
RDI: DRES Team [S. Piper (4/9/99), W. Cutchin (4/12/99)]; G. Kramer (4/12/99); M.Morrow (4/13/99)
J.Rowell:806W:CM#2:(703)305-5564:7509C:RAB1

Attachment 1: Pyridate Residue File for Acute and Chronic DEEM™ Analyses.

Filename: C:\MyFiles\DEEM\128834.r96

Chemical name: Pyridate

RfD(Chronic): .11 mg/kg bw/day NOEL(Chronic): 10.8 mg/kg bw/day

RfD(Acute): .2 mg/kg bw/day NOEL(Acute): 20 mg/kg bw/day

Date created/last modified: 04-02-1999/15:08:54/8

Program ver. 6.73

Comment: Pyridate on mint - G. Kramer. **The FQPA Safety Factor was removed, therefore the PAD and RfD are the same.**

Food Crop			RESIDUE	RDF	Adj. Factors		Comment
Code	Grp	Food Name	(ppm)	#	#1	#2	
259	6C	Beans-dry-garbanzo/chick pea	0.100000	0	1.000	1.000	
170	5A	Cabbage-green and red	0.030000	0	1.000	1.000	
383	5B	Cabbage-savoy	0.030000	0	1.000	1.000	
267	15	Corn grain-bran	0.030000	0	1.000	1.000	
266	15	Corn grain-endosperm	0.030000	0	1.000	1.000	
289	15	Corn grain-oil	0.030000	0	1.000	1.000	
268	15	Corn grain/sugar/hfcs	0.030000	0	1.500	1.000	
388	15	Corn grain/sugar-molasses	0.030000	0	1.500	1.000	
403	O	Peanuts-butter	0.030000	0	1.890	1.000	
940	O	Peanuts-hulled	0.030000	0	1.000	1.000	
293	O	Peanuts-oil	0.030000	0	1.000	1.000	

Attachment 2: Pyridate Acute DEEM™ Analysis (J. Rowell, 02-APR-1999).

U.S. Environmental Protection Agency
 DEEM ACUTE analysis for PYRIDATE
 Residue file: 128834.r96
 Analysis Date: 04-06-1999/11:15:24
 Acute Reference Dose (aRfD) = 0.200000 mg/kg body-wt/day
 NOEL (Acute) = 20.000000 mg/kg body-wt/day
 Run Comment: Pyridate on mint - G. Kramer. **The FQPA Safety Factor was removed, therefore the PAD and RfD are the same.**

Ver. 6.73

(1989-92 data)

Adjustment factor #2 NOT used.

Residue file dated: 04-05-1999/11:17:51/8

Summary calculations:

	95th Percentile		99th Percentile		99.9th Percentile	
	Exposure	% aRfD	Exposure	% aRfD	Exposure	% aRfD
U.S. pop - all seasons:	0.000139	0.07	0.000244	0.12	0.000406	0.20
Hispanics:	0.000149	0.07	0.000250	0.12	0.000421	0.21
Non-hispanic whites:	0.000135	0.07	0.000235	0.12	0.000409	0.20
Non-hispanic blacks:	0.000159	0.08	0.000278	0.14	0.000379	0.19
Non-hispanic other:	0.000128	0.06	0.000250	0.13	0.000594	0.30
All infants (<1 year):	0.000277	0.14	0.000411	0.21	0.000657	0.33
Nursing infants (<1 year):	0.000095	0.05	0.000142	0.07	0.000186	0.09
Non-nursing infants (<1 yr):	0.000278	0.14	0.000442	0.22	0.000682	0.34
Children (1-6 years):	0.000268	0.13	0.000391	0.20	0.000725	0.36
Children (7-12 years):	0.000179	0.09	0.000249	0.12	0.000378	0.19
Females (13+/preg/not nsg):	0.000082	0.04	0.000121	0.06	0.000200	0.10
Females (13+/nursing):	0.000124	0.06	0.000177	0.09	0.000194	0.10
Females (13-19 yrs/np/nn):	0.000109	0.05	0.000147	0.07	0.000320	0.16
Females (20+ years/np/nn):	0.000081	0.04	0.000130	0.07	0.000238	0.12
Females (13-50 years):	0.000092	0.05	0.000137	0.07	0.000268	0.13
Males (13-19 years):	0.000134	0.07	0.000208	0.10	0.000307	0.15
Males (20+ years):	0.000085	0.04	0.000132	0.07	0.000211	0.11
Seniors (55+):	0.000074	0.04	0.000117	0.06	0.000212	0.11

Attachment 3: Pyridate Chronic DEEM™ Analysis (J. Rowell, 02-APR-1999).

U.S. Environmental Protection Agency Ver. 6.74
 DEEM Chronic analysis for PYRIDATE (1989-92 data)
 Residue file name: C:\MyFiles\DEEM\128834.r96 Adjustment factor #2 NOT used.
 Analysis Date 04-02-1999/15:10:57 Residue file dated: 04-02-1999/15:08:54/8
 Reference dose (RfD, CHRONIC) = .11 mg/kg bw/day
 COMMENT 1: Pyridate on mint - G. Kramer. The FQPA Safety Factor was removed,
 therefore the PAD and RfD are the same.

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Total exposure by population subgroup

Population Subgroup	Total Exposure	
	mg/kg body wt/day	Percent of Rfd
U.S. Population (total)	0.000044	0.0%
U.S. Population (spring season)	0.000043	0.0%
U.S. Population (summer season)	0.000045	0.0%
U.S. Population (autumn season)	0.000045	0.0%
U.S. Population (winter season)	0.000042	0.0%
Northeast region	0.000040	0.0%
Midwest region	0.000045	0.0%
Southern region	0.000045	0.0%
Western region	0.000043	0.0%
Hispanics	0.000044	0.0%
Non-hispanic whites	0.000043	0.0%
Non-hispanic blacks	0.000050	0.0%
Non-hisp/non-white/non-black)	0.000041	0.0%
All infants (< 1 year)	0.000092	0.1%
Nursing infants	0.000023	0.0%
Non-nursing infants	0.000121	0.1%
Children 1-6 yrs	0.000105	0.1%
Children 7-12 yrs	0.000076	0.1%
Females 13-19(not preg or nursing)	0.000043	0.0%
Females 20+ (not preg or nursing)	0.000028	0.0%
Females 13-50 yrs	0.000032	0.0%
Females 13+ (preg/not nursing)	0.000031	0.0%
Females 13+ (nursing)	0.000036	0.0%
Males 13-19 yrs	0.000054	0.0%
Males 20+ yrs	0.000032	0.0%
Seniors 55+	0.000026	0.0%
Pacific Region	0.000041	0.0%



13544

R109035

Chemical: Pyridate

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